Table-S1 (Chemical structures of compounds with names and target activity)

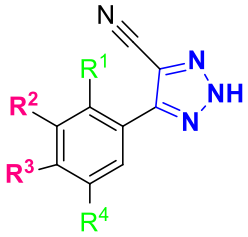
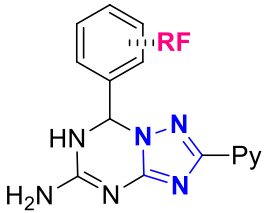
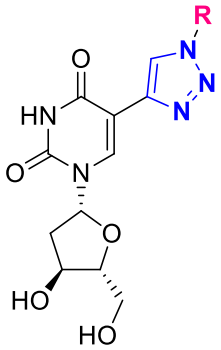
Chemical structure	No. of compounds / derivatives	Target/Activity	Reference
Anticancer Triazoles			
4-aryl-5-cyano-2H-1,2,3-triazole (1-2) 	**1**: R^1 = H; R^2 = H; R^3 = F; R^4 = H **2**: R^1 = H; R^2 = F; R^3= Pho; R^4 = H	Breast cancer cell line (MDA-MB-453)	(Cheng, Li et al. 2007)
7-aryl-2-pyridyl-6,7-dihydro[1,2,4]triazolo[1,5-a][1,3,5]triazin-5-amine (3-14) 	Py = 3-Py: RF = **3**: 2-F; **4**: 2-CF_3, **5**: 3-F, **6**: 3-CF_3, **7**: 4-F, **8**: 4-CF_3 Py = 4-Py, **9**: 2-F; **10**: 2-CF_3, **11**: 3-F, **12**: 3-CF_3, **13**: 4-F,**14**: 4-CF_3	Breast cancer cell line (MDA-MB-231), Colon cancer cell line (HT-29) And Lung cancer cell line A549,	(Dolzhenko, Tan et al. 2008)
1-((2R,4S,5R)-4-hydroxy-5-(hydroxymethyl)tetrahydrofuran-2-yl)-5-(1-Aryl-1H-1,2,3-triazol-4-yl)pyrimidine-2,4(1H,3H)-dione (15-19) 	**15**: R = -$CH_2CH_2CF_2CF_3$ **16**: R = -$CH_2CH_2(CF_2)_3CF_3$ **17**: R = -$CH_2CH_2(CF_2)_5CF_3$ **18**: R = -$CH_2CH_2(CF_2)_7CF_3$ **19**: R = -$CH_2CH_2(CF_2)_9CF_3$	Human breast cancer cell line (MDA-MB-231), Renal cancer cell line (ACHN), and Prostate cancer cell line (PC-3)	Park, Yang et al. 2010)

Table-S1 (Chemical structures of compounds with names and target activity)

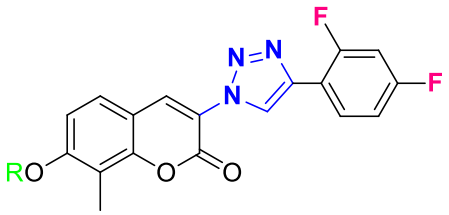
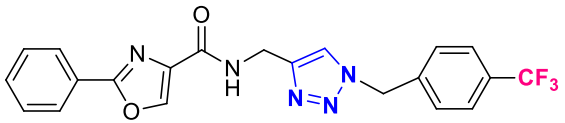
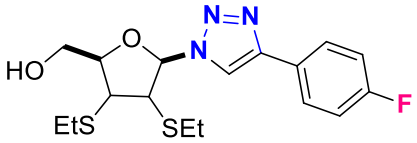
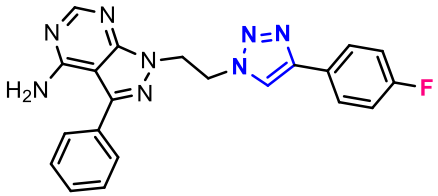
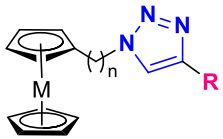
3-(4-(2,4-difluorophenyl)-1H-1,2,3-triazol-1-yl)-7-(Arylperoxy)-8-methyl-2H-chromen-2-one (20-21) 	**20**: R = OH **21**: R= OAc	Two breast cancer cell lines (MCF-7, SKBr-3)	(Peterson and Blagg 2010)
2-phenyl-N-((1-(4-(trifluoromethyl)benzyl)-1H-1,2,3-triazol-4-yl)methyl)oxazole-4-carboxamide (22) 	**22**	Breast cancer cell line (MCF-7), human leukemia cell line (HL-60), and Melanoma cell line (MDA-MB-435)	(Stefely, Palchaudhuri et al. 2010)
((2R,5R)-3,4-bis(ethylthio)-5-(4-(4-fluorophenyl)-1H-1,2,3-triazol-1-yl)tetrahydrofuran-2-yl)methanol (23) 	**23**	Human hepatocellular liver carcinoma cell line (HepG2)	(Yu, Wu et al. 2010)
1-(2-(4-(4-fluorophenyl)-1H-1,2,3-triazol-1-yl)ethyl)-3-phenyl-1H-pyrazolo[3,4-d]pyrimidin-4-amine (24) 	**24**	Breast cancer cell line (MDA-MB-361)	(Kumar, Ahmad et al. 2011)
Trifluoromethylated ferrocene triazole (25-28) 	**25**: M = Fe; n = 0; R = $(CF_3)_2OH$ **26**: M = Fe; n = 1; R = $(CF_3)_2OH$ **27**: M = Fe; n = 0; R = (CH_2SCF_3) **28**: M = Ru; n = 1; R = $(CF_3)_2OH$	Breast cancer cell line (MCF-7), colon cancer cell line (HT-29), and pancreas cancer cell line (PT-45)	(Maschke, Lieb et al. 2012)

Table-S1 (Chemical structures of compounds with names and target activity)

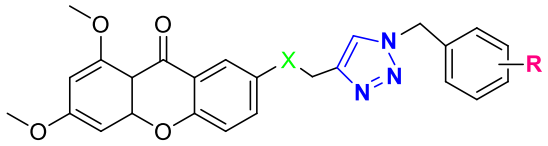
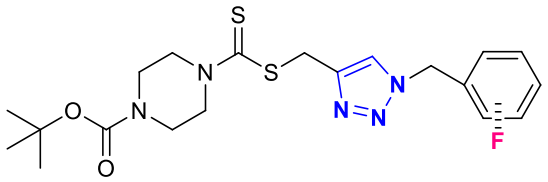
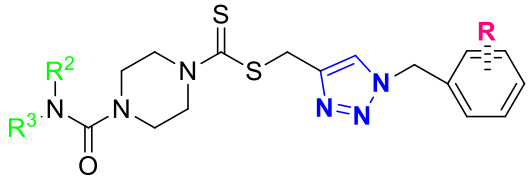
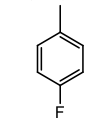
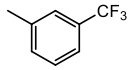
1,3-Disubstituted-1,2,3-triazole (29-34) 	X= NH; R = **29**: 2-F **30**: 3-F **31**: 4-F X= O; R = **32**: 2-F **33**: 3-F **34**: 4-F	Hepatoma carcinoma cell line (Bel-7402) and human cervical carcinoma cell line (HeLa)	(Zou, Zhao et al. 2012)
Tert-butyl4-((((1-(Fluorobenzyl)-1H-1,2,3-triazol-4 yl)methyl)thio)carbonothioyl)piperazine-1-carboxylate (35-36) 	**35**: *o*-F **36**: *m*-F	Human gastric cancer cell line (MGC-803) and Human breast cancer cell line (MCF-7)	(Duan, Ma et al. 2013)
1,2,3-triazole-dithiocarbamate-urea hybrids (37-38) 	$R^2R^3NH = (CH_3)_2CHNH$ **37**: R^1 = *o*-F **38**: R^1 = *p*-F	Human breast cancer cell line (MCF-7), Human gastric cancer cell line (MGC-803) and Hepatocellular carcinoma cell line (SMMC-7721), human esophageal cancer cell line (EC-9706)	(Duan, Zheng et al. 2013)
6-[(N1-aryl-1H-1,2,3-triazol-4-yl)methyl]-6H-indolo[2,3-b]quinoxa-l (39-43)	R = H, F; R^1= **39**:  **40**: 	Cervical cancer cell line (HeLa), prostate cancer cell line (DU-145), and lung cancer cell line (A-549)	(Avula, Komsani et al. 2012)

Table-S1 (Chemical structures of compounds with names and target activity)

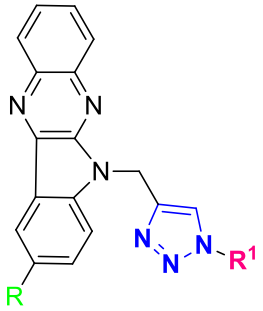
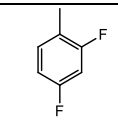
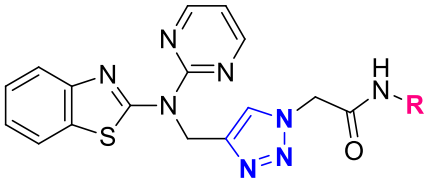
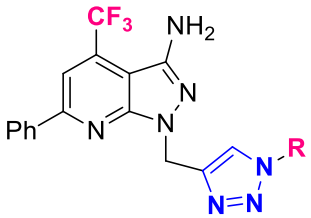
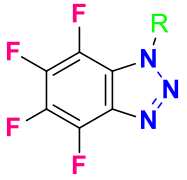
	**41**:  **42**:		
Novel triazole linked N-(pyrimidin-2-yl)benzo[d]thiazol-2-amine derivatives (43-50) 	**43**: R = CH_2-4-$OCF_3C_6H_4$ **44**: R = 4-$SCF_3C_6H_4$ **45**: R = 5(CF_3)-1,3,4-Thiazole **46**:R=4-F-benzothiazole morpholine **47**: R = 4-FC_6H_4 **48**: R = 2,4-FC_6H_3 **49**: R = 3-$CF_3C_6H_4$ **50**: R = (3-Cl-4-FC_6H_3)	Lung (A549), breast (MCF-7) and skin (A375) cancer cell lines	(Kumbhare, Dadmal et al. 2015)
1,2,3-triazole tagged pyrazolo[3,4-b]pyridine derivatives (51-56) 	**R=** **51**: C_6F_{13}-CH_2-CH_2- **52**: C_8F_{17}-CH_2-CH_2- **53**: 3-Cl, 4-FC_6H_3- **54**: 3-$CF_3C_6H_4$- **55**: 4-FC_6H_4- **56**: 3-F,4-BrC_6H_3-	Human monocytic leukemia (U937), human acute monocytic leukemia (THP-1), human promyelocytic leukemia (HL-60) and Mouse Me (B16-F10) cancer cell lines	(Kurumurthy, Veeraswamy et al. 2014)
4,5,6,7-tetrafluoro-1Aryl-benzo[d][1,2,3]triazole (57-60) 	**57**: R = H **58**: R = Cl **59**: R = H_3CO **60**: R = $(CH_3)_2N$	Hep2 (laryngeal epidermoid carcinoma) cancer cells	(Prima, Baev et al. 2017)

Table-S1 (Chemical structures of compounds with names and target activity)

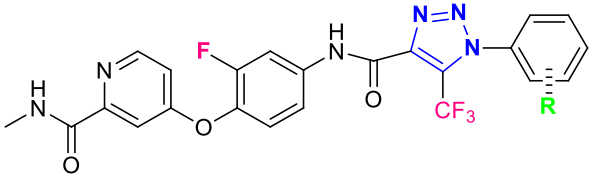
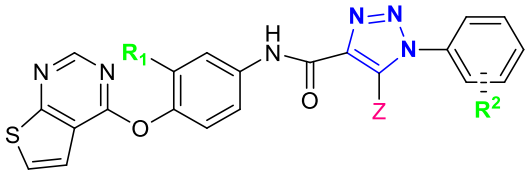
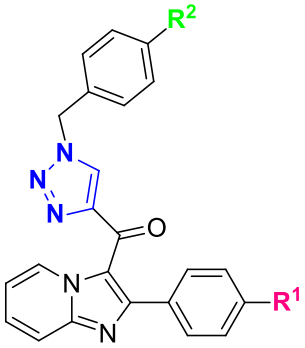
Thinopyrimidine-triazole conjugates (61-71)  	Z= CF_3; R = **61**: 2-CF_3; **62**: 4-Cl; **63**: 3-F-4-F **64**: 3-F-4-Cl, **65**: 4-Cl-3-CF_3 Z = CH_3 **66**: 4-F R^1 = H; Z = CF_3; R^2 = **67**: 2-CF_3 **68**: 4-Cl **69**: 3-F-4-F **70**: 2-F-4-F **71**: F-F-4-Cl	Human lung cancer cell line (A549), Human liver cancer cell line (HepG2) and Human breast cancer cell line (MCF-7).	(Wang, Xu et al. 2018)
(1-((Arylphenyl]-4-yl)methyl)-1H-1,2,3-triazol-4-yl)(2-(4-Arylphenyl)imidazo[1,2-a]pyridin3yl)methanone (72-74) 	R^1 = 4-flourophenyl; **72**: R^2 = OCH3 **73**: R^2 = H **74**: R^2 = Cl	Human lung cancer (A549), Human prostate (HCT-116), and Human breast cancer cell (MDA-MB-231)	(Sayeed, Vishnuvardhan et al. 2018)
(1aR,7aS,10aS,10bR,E)-5-((4-(3,5-bis(trifluoromethyl)phenyl)-1H-1,2,3-triazol-1-yl)methyl)-1a-methyl-8-methylene-2,3,6,7,7a,8,10a,10b-octahydrooxireno[2',3':9,10]cyclodeca[1,2-b]furan-9(1aH)-one (75)	**75**	M9 ENL1 cell, (AML#1, and AML#2)	(Janganati, Ponder et al. 2018)

Table-S1 (Chemical structures of compounds with names and target activity)

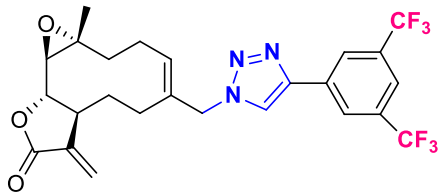
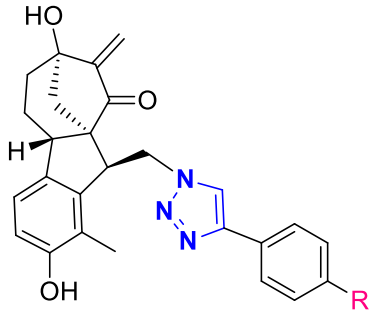
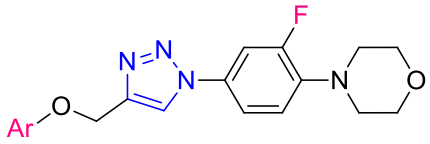
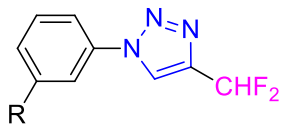
			
(4bS,7S,9aR,10S)-10-((4-(Aryl-phenyl)-1H-1,2,3-triazol-1-yl)methyl)-2,7-dihydroxy-1-methyl-8-methylene-5,6,7,8-tetrahydro-4bH-7,9a-methanobenzo[a]azulen-9(10H)-one (76-77) 	**76**: R = F **77**: R = CF_3	Human myeloid leukaemia (HL-60), human lung carcinoma (A549), human liver carcinoma (SMMC-7721), human colon carcinoma (SW480) and human breast carcinoma (MCF-7) cell lines	(Wu, Wu et al. 2018)
4-[3-fluoro-4-(morpholin-4-yl)]phenyl-1H-1,2,3-triazole (78-86) 	Ar = **78**: C_6H_5, **79**: 2-FC_6H_4, **80**: 4-FC_6H_4 **81**: 3MeC_6H_4, **82**: 3,5 diMeC_6H_3 **83**: 4$MeOC_6H_4$, **84**: 3ClC_6H_4 **85**: 3,5 diClC_6H_3, **86**: 4-BrC_6H_4	Breast cancer cell line (MCF-7) and cervical carcinoma cell line (HeLa)	(Narsimha, Nukala et al. 2020)
Anti-bacterial Triazoles			
1-(4-Arylphenyl)-1H-1,2,3-triazole-4-carbaldehyde (87-89) 	**87**: R= 4-Cl **88**: R= 4-Br **89**: R= 4-CH_3	Mycobacterium tuberculosis	(Costa, Boechat et al. 2006)

Table-S1 (Chemical structures of compounds with names and target activity)

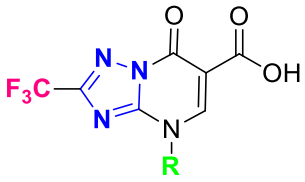
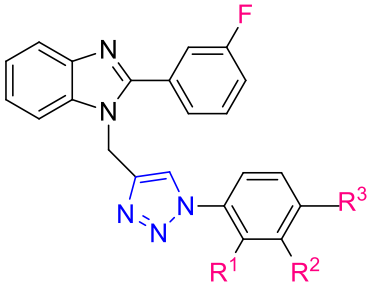
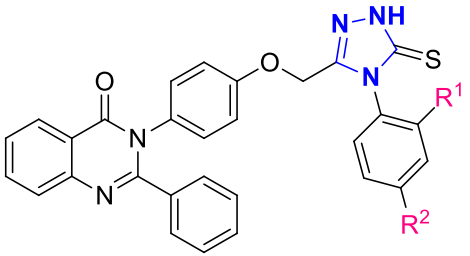
4-Aryl-7-oxo-2-(trifluoromethyl)-4,7-dihydro-1,2,4-triazolo[5,1-a]pyrimidine-6-carboxylic acid (90-94) 	**90**: R = CH_3 **91**: R = C_2H_5 **92**: R = $-CH_2CH=CH_2$ **93**: R = $-CH_2-C_6H_4(4\text{-}Br)$ **94**: R = $-CH_2-C_6H_4(4\text{-}NO_2)$	Mycobacterium tuberculosis	(Abdel-Rahman, El-Koussi et al. 2009)
2-(3-fluorophenyl)-1-((1-(Arylphenyl)-1H-1,2,3-triazol-4-yl)methyl)-1H-benzo[d]imidazole (95-100) 	**95**: R^1 = F; R^2 = F; R^3 = F **96**: R^1 = H; R^2= F; R^3 = F **97**: R^1 = H; R^2 = F; R^3 = H **98**: R^1 = F; R^2 = H; R^3 = Me **99**: R^1 = F; R^2 = H; R^3 = F **100**: R^1 = H; R^2 = H; R^3 = CF_3	(staphylococcus aureus, Pseudomonas aeruginosa) (Escherichia coli, Salmonella typhosa)	(Gill, Jadhav et al. 2008)
3-(4-((4-(Arylphenyl)-5-thioxo-4,5-dihydro-1H-1,2,4-triazol-3-yl)methoxy)phenyl)-2-phenylquinazolin-4(3H)-one (101-102) 	**101**: R^1 = H; R^2= F **102:** R^1 = F; R^2= F	Escherichia coli. Bacillus subtilis and Staphylococcus aureus.	(Havaldar, F. et al. 2008)

Table-S1 (Chemical structures of compounds with names and target activity)

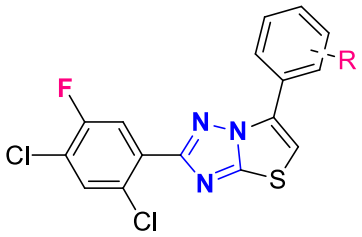
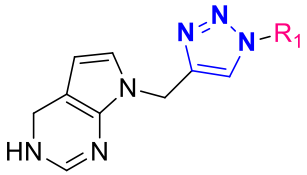
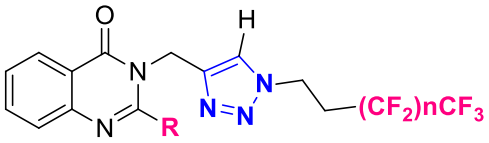
(2-(2,4-dichloro-5-fluorophenyl)-6-(aryl)thiazolo[3,2-b][1,2,4]triazole) (103-110) 	**103**: R = 4-OCH_3 **104**: R = 4-CH_3 **105**: R = 4-F **106**: R = 4-Cl **107**: R = 4-Br **108**: R = 4-NO_2 **109**: R = 2,4-Cl2 **110**: R = 2,4-Cl2-5-F	Escherichia coli, Staphylococcus aureus, Pseudomonas aeruginosa, Streptococcus pyogenes and Klebsiella pneumoniae	(Karthikeyan 2009)
7-((1-(Aryl)-1H-1,2,3-triazol-4-yl)methyl)-4,7-dihydro-3H-pyrrolo[2,3-d]pyrimidine (111-112) 	**111**: R^1 = *o*-F-Ph, **112**: R^1 = *p*-F-Ph	Mycobacterium tuberculosis	(Shiva Raju, AnkiReddy et al. 2019)
2-Aryl-3-((1-(Aryl)-1H-1,2,3-triazol-4-yl)methyl) quinazolin-4(3H)-one (113-118) 	**113**: R = C_6H_5; n = 5 **114**: R = C_6H_5; n = 7 **115**: R = 2,6-$C_6H_3F_2$; n = 5 **116**: R = 2,6-$C_6H_3F_2$; n = 7 **117**: R = CF_3; n = 5 **118**: R = CF_3; n = 7	Gram positive bacteria (Bacillus subtilis, Staphylococcus aureus, Staphylococcus epidermidis) and gram negative bacteria (Pseudomonas aeruginosa, Escherichia coli)	(Mani Chandrika, Yakaiah et al. 2010)

Table-S1 (Chemical structures of compounds with names and target activity)

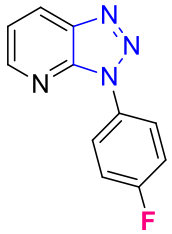
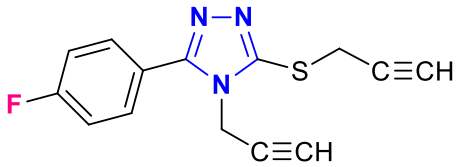
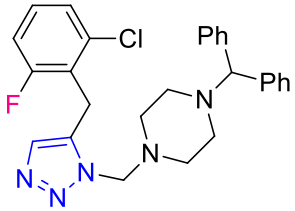
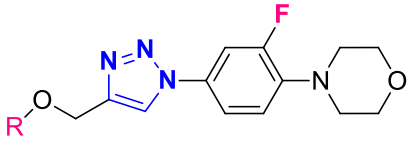
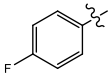
3-(4-fluorophenyl)-3H-[1,2,3]triazolo[4,5-b]pyridine (119) 	**119**	Bacillus subtilis and Escherichia coli	(Marepu, Yeturu et al. 2018)
3-(4-fluorophenyl)-4-(prop-2-yn-1-yl)-5-(prop-2-yn-1-ylthio)-4H-1,2,4-triazole (120) 	**120**	Bacillus subtilis, Streptococcus pneumonia, Staphylococcus aureus, Escherichia coli, Pseudomonas aeruginosa, and Klebsiella pneumonia	(Rezki, Mayaba et al. 2016)
1-benzhydryl-4-((5-(2-chloro-6-fluorobenzyl)-1H-1,2,3-triazol-1-yl)methyl)piperazine (121) 	**121**	Staphylococcus aureus and Escherichia coli.	(Govindaiah, Sreenivasa et al. 2018)
4-(2-fluoro-4-(4-([2-Aryloxy]methyl)-1H-1,2,3-triazol-1-yl)phenyl)morpholine (122-123) 	**122**: R =  **123**: R = 	Bacillus subtilis, Staphylococcus aureus, and Staphylococcus epidermidis	(Narsimha, Nukala et al. 2020)

Table-S1 (Chemical structures of compounds with names and target activity)

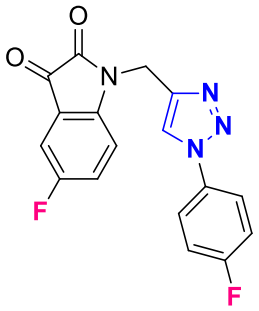
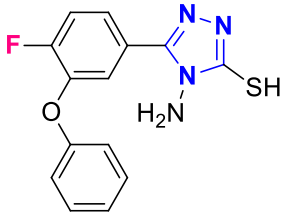
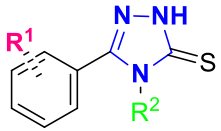
5-fluoro-1-((1-(4-fluorophenyl)-1H-1,2,3-triazol-4-yl)methyl)indoline-2,3-dione (124) 	**124**	Staphylococcus epidermidis, Bacillus subtilis, Escherichia coli, and Pseudomonas aeruginosa	(Deswal, Naveen et al. 2020)
4-amino-5-(4-fluoro-3-phenoxyphenyl)-4H-1,2,4-triazole-3-thiol (125) 	**125**	Mycobacterium tuberculosis	(Venugopala, Kandeel et al. 2020)
3-(Aryl-phenyl)-4-(Aryl)-1H-1,2,4-triazole-5(4H)-thione (126-131) 	R^1 = *o*-F, R^2 = **126**: 1-naph **127**: *m*-tol **128**: *p*-tol R^1 = *o*-F, *m*-F **129**: R^2 = 1-naph **130**: R^2 = *m*-tol **131**: R^2 = *p*-tol	Staphylococcus aureus,	(Kosikowska, Wujec et al. 2020)

Table-S1 (Chemical structures of compounds with names and target activity)

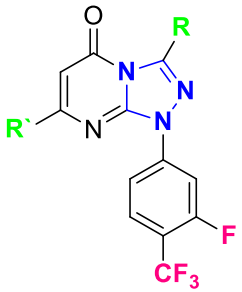
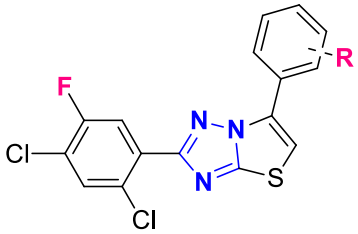
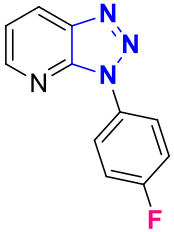
3-Aryl-1-(3-fluoro-4-methylphenyl)-7Aryl-[1,2,4]triazolo[4,3-a]pyrimidin-5(1H)-one (132-135) 	R/R*: **132**: a = $COCH_3/CH_3$ **133**: b = COOEt /CH_3 **134**: c = $COCH_3$/Ph **135**: d = COOE/Ph	Bacillus subtilis and Escherichia coli	(Muhammad, Farghaly et al. 2021)
Antifungal Triazoles			
(2-(2,4-dichloro-5-fluorophenyl)-6-(aryl)thiazolo[3,2-b][1,2,4]triazole) (136-143) 	**136**: R = 4-OCH_3 **137**: R = 4-CH_3 **138**: R = 4-F **139**: R = 4-Cl **140**: R = 4-Br **141**: R = 4-NO_2 **142**: R = 2,4-Cl_2 **143**: R = 2,4-Cl_2-5-F	Aspergillus niger, A. fumigstus, Candida albicansa, Penicillium marneffei, and Trichophyton mentagrophytes	(Karthikeyan 2009)
3-(4-fluorophenyl)-3H-[1,2,3]triazolo[4,5-b]pyridine (144) 	**144**	f.sp.Lycopersici. Fusarium oxysporium and Fusarium ricini	(Marepu, Yeturu et al. 2018)
2-Aryl-3-((1-(Aryl)-1H-1,2,3-triazol-4-yl)methyl) quinazolin-4(3H)-one (145-150)	**145**: R = C_6H_5; n = 5 **146**: R = C_6H_5; n = 7 **147**: R = 2,6-$C_6H_3F_2$; n = 5 **148**: R = 2,6-$C_6H_3F_2$; n = 7 **149**: R = CF_3; n = 5	Candida albicans, Saccharomyces cerevisiae and filamentous fungal culture like	(Mani Chandrika, Yakaiah et al. 2010)

Table-S1 (Chemical structures of compounds with names and target activity)

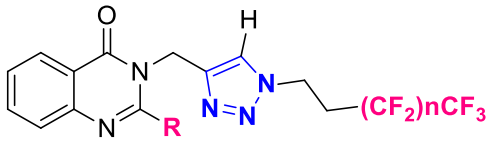
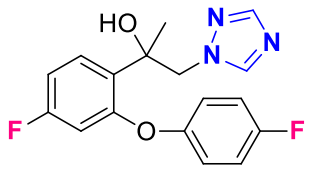
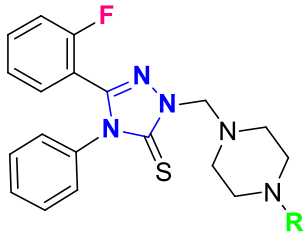
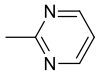
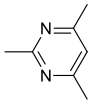
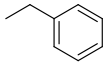
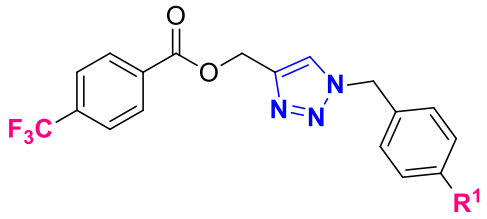
	**150**: R = CF_3; n = 7	Rhizopus oryzae, Aspergillus flavus and Candida rugosa	
2-(4-fluoro-2-(4-fluorophenoxy)phenyl)-1-(1H-1,2,4-triazol-1-yl)propan-2-ol (151) 	**151**	Gibberella zeae, Alternaria solani, Fusarium oxysporium, Physalospora pirco	(Yang, Zhai et al. 2015)
N-(1-((4-substituted piper112 azin-1-yl)methyl)-3-methyl-5-thioxo-1H-1,2,4-triazol-4(5H)-yl)-2/3-fluorobenzamide (152-155) 	R= **152**:  **153**:  **154**:  **155**: 	Cercospora arachidicola, Physalospora piricola, Rhizoctonia cerealis	(Zhang, Wang et al. 2016)
(1-(Aryl)-1H-1,2,3-triazol-4-yl)methyl 4-(trifluoromethyl)benzoate (156-157) 	**156**: R^1 = 4-F **157**: R^1 = 4-F-C_6H_5	Aspergillus niger and Candida. albicans	(Deswal, Tittal et al. 2019)

Table-S1 (Chemical structures of compounds with names and target activity)

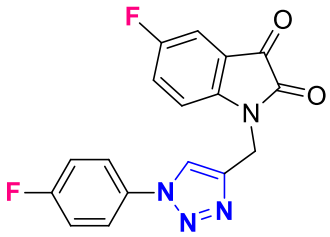
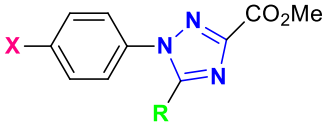
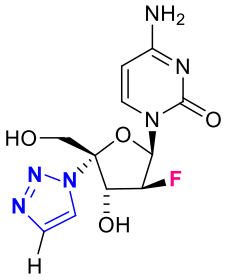
5-fluoro-1-((1-(4-fluorophenyl)-1H-1,2,3-triazol-4-yl)methyl)indoline-2,3-dione (158) 	**158**	Aspergillus niger, And Candida albicans,	(Deswal, Naveen et al. 2020)
Antiproliferative Triazoles			
Trifluoromethane-3,5-disubstituted 1,2,4-triazoles (159-162) 	**159**: X = p-F; R = Me **160**: X = p-CF_3; R = Me **161**: X = m-CF_3; R = Me **162**: X = p-F; R = Et	NPC-TW01 And T-cell leukemia cell	(Wang, Tseng et al. 2011)
Antiviral Triazoles			
4-amino-1-((2R,3S,4R,5R)-3-fluoro-4-hydroxy-5-(hydroxymethyl)-5-(1H-1,2,3-triazol-1yl)tetrahydrofuran-2-yl)pyrimidin-2(1H)-one (163) 	**163**	293T cell line	(Wu, Yu et al. 2013)

Table-S1 (Chemical structures of compounds with names and target activity)

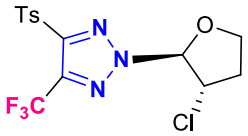
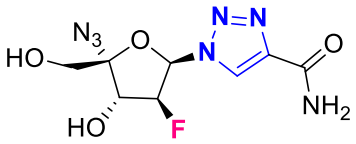
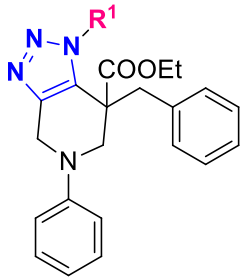
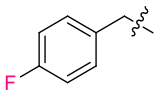
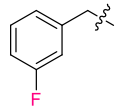
2-(trans-3-Chlorotetrahydrofuran-2-yl)-4-tosyl-5-(trifluoromethyl)-2H1,2,3-triazole (164) 	**164**	HSV-1, HSV-2 and HAdV5	(Biliavska, I., et al. 2017)
1-((2R,3S,4R,5R)-5-azido-3-fluoro-4-hydroxy-5-(hydroxymethyl) tetrahydrofuran-2-yl)-1H-1,2,3-triazole-4-carboxamide (165) 	**165**	Wild-type HBV	(Liu, Peng et al. 2018)
Ethyl7-benzyl-1-(Aryl)-5-phenyl-4,5,6,7-tetrahydro-1H-[1,2,3]triazolo[4,5-c]pyridine-7-carboxylate (166-167) 	**166**: R^1 =  **167**: R^1 = 	Coronavirus (229E)	(Karypidou, Ribone et al. 2018)
Antimicrobial Triazoles			
2-(3-fluorophenyl)-1-((1-(Arylphenyl)-1H-1,2,3-triazol-4-yl)methyl)-1H-benzo[d]imidazole(168-173)	**168**: R^1 = F; R^2 = F; R^3 = F **169**: R^1 = H; R^2 = F; R^3 = F **170**: R^1 = H; R^2 = F; R^3 = H **171**: R^1 = F; R^2 = H; R^3 = Me **172**: R^1 = F; R^2 = H; R^3 = F **173**: R^1 = H; R^2 = H; R^3 = CF_3	Staphylococcus aureus, Pseudomonas aeruginosa, Escherichia coli and Salmonella typhosa.	(Gill, Jadhav et al. 2008)

Table-S1 (Chemical structures of compounds with names and target activity)

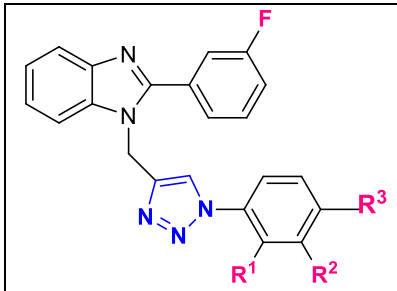
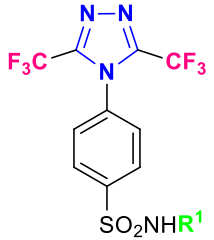
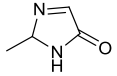
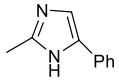
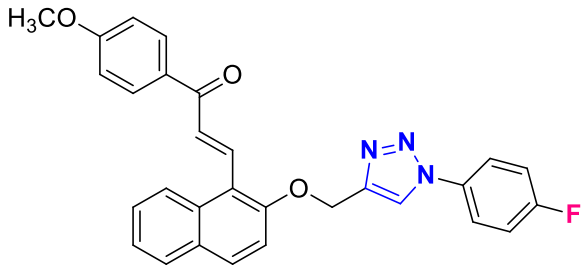
			
N-di(trifluoroacetyl)sulfonamide (174-177) 	**174**: R^1 = H, **175**: R^1 = CH_3CO, **176**: R^1 =  **177**: R^1 = 	Staphylococcus aureus, Escherichia coli, Aspergillus niger and Candida albicans	(Faidallah, Khan et al. 2011)
(E)-3-(2-((1-(4-fluorophenyl)-1H-1,2,3-triazol-4-yl)methoxy) naphthalen-1-yl)-1-(4-methoxyphenyl)prop-2-en-1-one (178) 	**178**	Staphylococcus epidermidis, Bacillus subtilis, Escherichia. coli, Pseudomonas. aeruginosa, Aspergillus niger and Candida albicans	(Yadav, Lal et al. 2018)
Herbicidal Triazoles			

Table-S1 (Chemical structures of compounds with names and target activity)

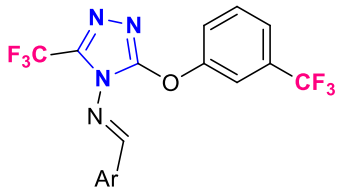
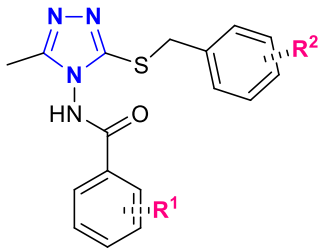
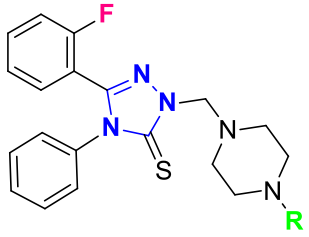
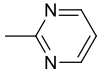
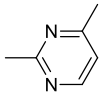
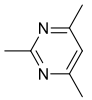
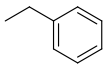
(E)-N-(argiomethylene)-3-(trifluoromethyl)-5-(3-(trifluoromethyl)phenoxy)-4H-1,2,4-triazol-4-amine (179) 	**179**	Brassica campestris, and Echinochloa crus-galli	(Zhang and Shi 2014)
Aryl-N-(3-((Arylbenzyl)thio)-4H-1,2,4-triazol-4-yl)benzamide (180-185) 	R^1 = *o*-F; **180**: R^2 = *o*-F **181**: R^2 = *m*-F **182**: R^2 = *p*-F R^1 = *m*-F; **183**: R^2 = *o*-F **184**: R^2 = *m*-F **185**: R^2 = *p*-F	Brassica campestris, and Echinochloa crusgalli	(Liu, Weng et al. 2013)
N-(1-((4-substituted piper112 azin-1-yl)methyl)-3-methyl-5-thioxo-1H-1,2,4-triazol-4(5H)-yl)-2/3-fluorobenzamide (186-189) 	R = **186**:  **187**:  **188**:  **189**: 	Brassica campestris and Echinochloa crus-galli	(Zhang, Wang et al. 2016)
Inhibitory Triazoles			

Table-S1 (Chemical structures of compounds with names and target activity)

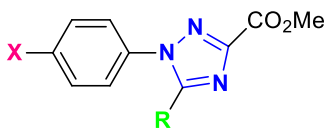
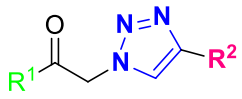
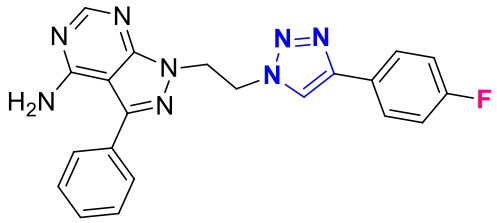
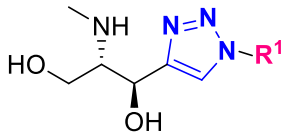
Trifluoromethane-3,5-disubstituted 1,2,4-triazoles (190-193) 	**190**: X = *p*-F; R = Me **191**: X = *p*-CF_3; R = Me **192**: X = *m*-CF_3; R = Me **193**: X = *p*-F; R = Et	Lung carcinoma, Nasopharyngeal, And T-cell leukemia (Jurkat) cells	(Wang, Tseng et al. 2011)
2-(4-(Aryl)-1H-1,2,3-triazol-1-yl)-1-phenylethanone (194-197) 	**194**; $R^1 = C_6H_5$; R^2= 4-F-3-$CH_3C_6H_3$ **195**:R^1=4-$CH_3C_6H_4$;R^2=4-F-3-$CH_3C_6H_3$ **196**: R^1 = 4-ClC_6H_4; R^2= 4-FC_6H_4 **197**: R^1 = 4-BrC_6H_4; R^2= 4-FC_6H_4	Src kinase	(Kumar, Reddy et al. 2011)
1-(2-(4-(4-fluorophenyl)-1H-1,2,3-triazol-1-yl)ethyl)-3-phenyl-1H-pyrazolo[3,4-d]pyrimidin-4amine (198) 	**198**	Src kinase	(Kumar, Ahmad et al. 2011)
Tert-butyl((1S,2S)-1,3-dihydroxy-1-(1-Aryl-1H-1,2,3-triazol-4-yl)propan-2-yl) (199-201) 	**199**: R^1; $C_{11}H_{22}CF_3$ 61% **200**: R^1; $C_{10}H_{20}CF_2CF_3$ 63% **201**: R^1; $C_9H_{18}CF_2CF_2CF_3$ 69%	SPHK1 and SPHK2	(Escudero-Casao, Cardona et al. 2018)
2,3,5,6-tetrafluoro-4-(4-Aryl-1H-1,2,3-triazol-1-yl)benzenesulfinamide (202-209)	R = **202**: C_6H_5 **203**: $BrCH_2CH_2$ **204**: MeOOC **205**: *p*-Me-C_6H_4 **206**: $Cl(CH_2)_3$	Isoform hCA I, II, IX and XII	(Pala, Micheletto et al. 2014)

Table-S1 (Chemical structures of compounds with names and target activity)

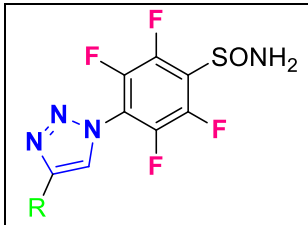
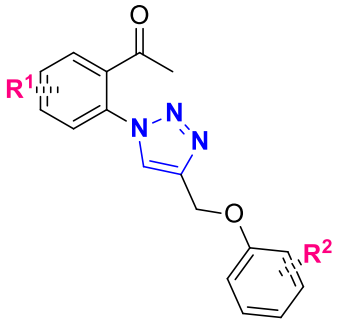
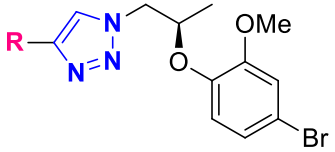
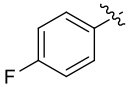
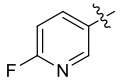
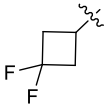
	**207**: $Cl(CH_2)_4$ **208**: $HO\text{-}CH_2$ **209**: $H2N\text{-}CH_2$		
2-(4-((Aryl-phenoxy)methyl)-Aryl-1,2,3-triazol-1-yl)benzamide(210-216) 	R^1= H; R^2 = **210**: 4-F **211**: $4\text{-}CF_3$ **212**: $3\text{-}CF_3$ **213**: 3,4-2F R^1= 4,5-2F; R^2 = **214**: 4-F **215**: $4\text{-}CF_3$ **216**: 3,4-2F	Hdhodh	(Lu, Cai et al. 2018)
(R)-1-(2-(4-bromo-2-methoxyphenoxy)propyl)-4-(Aryll)-4-yl)-1H-1,2,3-triazole (217-218) 	**217**: R = $2\text{-}CF_3C_6H_4$ **218**: R = $4\text{-}FC_6H_4$	α-glucosidase enzyme	(Avula, Khan et al. 2018)
(S)-N-(3-(3-chloro-4-(trifluoromethoxy)phenyl)-1-((1-cyanocyclopropyl)amino)-1-oxopropan-2-yl)-2-(Aryl)-2H-1,2,3triazole-4-carboxamide (219-221)	R= **219**:  **220**:  **221**: 	Human cathepsin L (hCatL)	(Giroud, Kuhn et al. 2018)

Table-S1 (Chemical structures of compounds with names and target activity)

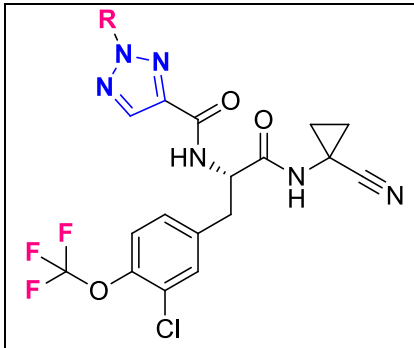
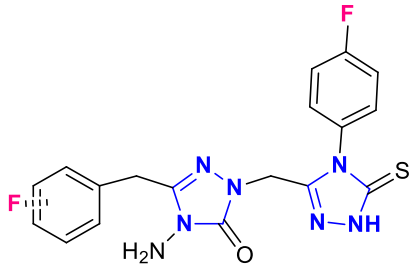
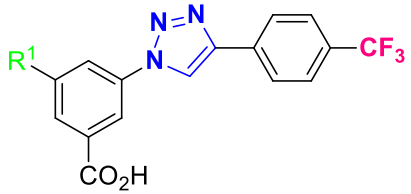
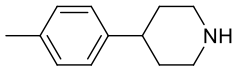
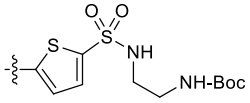
			
Antioxidant Triazoles			
4-amino-3-(3/4-fluorobenzyl)-1-((4-(4-fluorophenyl)-5-thioxo-4,5-dihydro-1H-1,2,4-triazol-3-yl)methyl)-1H-1,2,4-triazol-5(4H)-one (222-223) 	**222**: *m*-F **223**: *p*-F	Jack bean urease	(Bekirican, O. et al. 2016)
Antagonistic Triazoles			
3-(Aryl)-5-(4-(4-(trifluoromethyl)phenyl)-1H-1,2,3-triazol-1-yl)benzoic acid (224-227) 	R^1 = **224**: -OH **225**:  **226**: 	P2YR14R	(Yu, Ciancetta et al. 2018)

Table-S1 (Chemical structures of compounds with names and target activity)

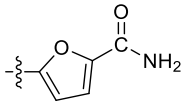
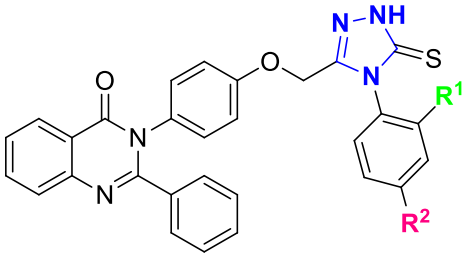
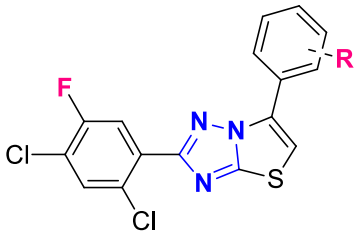
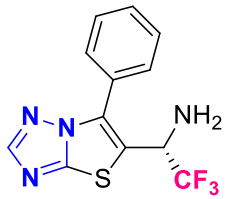
	227: 		
Antimalarial Triazoles			
3-(4-((4-(Arylphenyl)-5-thioxo-4,5-dihydro-1H-1,2,4-triazol-3-yl)methoxy)phenyl)-2-phenylquinazolin-4(3H)-one (228-229) 	**228**: R^1 = H; R^2 = F **229**: R^1 = F; R^2 = F	Plasmodium falciparum	(Havaldar, F. et al. 2008)
Anti-inflammatory Triazoles			
(2-(2,4-dichloro-5-fluorophenyl)-6-(aryl)thiazolo[3,2-b][1,2,4]triazole) (230 – 237) 	R = **230**: 4-OCH_3 **231**: 4-CH_3 **232**: 4-F **233**: 4-Cl **234**: 4-Br **235**: 4-NO_2 **236**: 2,4-Cl_2 **237**: 2,4-Cl_2-5-F	*Escherichia coli*, *Staphylococcus aureus*, *Aspergillus niger* and *Candida albicans*	(Karthikeyan 2009)
(R)-2,2,2-trifluoro-1-(6-phenylthiazolo[3,2-b][1,2,4]triazol-5-yl)ethanamine (238)	**238**	TT-TFM	(Tu, Yu et al. 2018)

Table-S1 (Chemical structures of compounds with names and target activity)

"<Antimalarial activity-1.pdf>."
"<Antioxidant activity-1.pdf>."
Abdel-Rahman, H. M., N. A. El-Koussi and H. Y. Hassan (2009). "Fluorinated 1,2,4-Triazolo[1,5-a]pyrimidine-6-carboxylic acid derivatives as antimycobacterial agents." Arch Pharm (Weinheim) **342**(2): 94-99.
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Table-S1 (Chemical structures of compounds with names and target activity)

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